Gary James Connett

List of Publications by Year
in descending order

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103	5,483	172457	82547
papers	citations	29 h-index	72 g-index

105	105	105	6645
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Projecting the impact of triple CFTR modulator therapy on intravenous antibiotic requirements in cystic fibrosis using patient registry data combined with treatment effects from randomised trials. Thorax, 2022, 77, 873-881.	5.6	11
2	Structured transition is associated with improved outcomes in diabetes. Practical Diabetes, 2022, 39, 18.	0.3	1
3	Evaluation of the impact of shielding to avoid COVID-19 infection on respiratory symptoms in children with severe asthma. Archives of Disease in Childhood, 2021, 106, e23-e23.	1.9	3
4	The impact of plasma 25â€hydroxyvitamin D on pulmonary function and exercise physiology in cystic fibrosis: A multicentre retrospective study. Journal of Human Nutrition and Dietetics, 2021, , .	2.5	2
5	Meconium Ileus due to GUCY2C gene mutations in three unrelated South Indian families. Journal of Cystic Fibrosis, 2021, 20, e84-e86.	0.7	1
6	The implications of dysglycaemia on aerobic exercise and ventilatory function in cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, 427-433.	0.7	8
7	Cephalosporin nitric oxide-donor prodrug DEA-C3D disperses biofilms formed by clinical cystic fibrosis isolates of Pseudomonas aeruginosa. Journal of Antimicrobial Chemotherapy, 2020, 75, 117-125.	3.0	35
8	A public health emergency among young people. Lancet Respiratory Medicine,the, 2020, 8, 231-233.	10.7	6
9	The impact of plasma 25-hydroxyvitamin d on lung function in patients with cystic fibrosis. Clinical Nutrition ESPEN, 2020, 40, 537-538.	1.2	0
10	The ERS approach to e-cigarettes is entirely rational. European Respiratory Journal, 2020, 55, 2000413.	6.7	2
11	Circulating biomarkers of antioxidant status and oxidative stress in people with cystic fibrosis: A systematic review and meta-analysis. Redox Biology, 2020, 32, 101436.	9.0	35
12	Central and obstructive apnoea-hypopnea indices pre and post growth hormone commencement in children with Prader-Willis syndrome following introduction of a screening programme. , 2020, , .		0
13	Lumacaftor-ivacaftor in the treatment of cystic fibrosis: design, development and place in therapy . Drug Design, Development and Therapy, 2019, Volume 13, 2405-2412.	4.3	35
14	Efficacy and safety of the elexacaftor plus tezacaftor plus ivacaftor combination regimen in people with cystic fibrosis homozygous for the F508del mutation: a double-blind, randomised, phase 3 trial. Lancet, The, 2019, 394, 1940-1948.	13.7	804
15	P415 Medicine possession ratios for ivacaftor prescriptions data in children and adults with cystic fibrosis attending a UK regional centre. Journal of Cystic Fibrosis, 2019, 18, S174-S175.	0.7	0
16	P209 Use of lumacaftor/ivacaftor as rescue therapy and stabilisation treatment for severe lung disease in children with cystic fibrosis. Journal of Cystic Fibrosis, 2019, 18, S116.	0.7	0
17	Determining the reasons for poorly controlled asthma in an adolescent. BMJ: British Medical Journal, 2019, 364, l75.	2.3	2
18	Reply to Askew and Green. Journal of Applied Physiology, 2019, 126, 512-512.	2.5	0

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19	Reply to Cooper. Journal of Applied Physiology, 2019, 126, 265-265.	2.5	0
20	Psychometric evaluation of a patient-reported outcome measure in pancreatic exocrine insufficiency (PEI). Pancreatology, 2019, 19, 182-190.	1.1	32
21	Airway Microbiome and Development of Bronchopulmonary Dysplasia in Preterm Infants: A Systematic Review. Journal of Pediatrics, 2019, 204, 126-133.e2.	1.8	81
22	Dysfunctional Breathing in Children and Adults With Asthma. Frontiers in Pediatrics, 2018, 6, 406.	1.9	34
23	Ready, Steady, Go â€“ Achieving successful transition in cystic fibrosis. Paediatric Respiratory Reviews, 2018, 27, 13-15.	1.8	11
24	The use of lumacaftor/ivacaftor to treat acute deterioration in paediatric cystic fibrosis. Paediatric Respiratory Reviews, 2018, 27, 16-17.	1.8	3
25	Evaluation of a regionally based preceptorship programme for newly qualified neonatal nurses. Journal of Neonatal Nursing, 2018, 24, 225-228.	0.7	2
26	Cardiopulmonary exercise testing with supramaximal verification produces a safe and valid assessment of V̇I‡<scp>o</scp>_{2max} in people with cystic fibrosis: a retrospective analysis. Journal of Applied Physiology, 2018, 125, 1277-1283.	2.5	27
27	Cephalosporin-3â€²-Diazeniumdiolate NO Donor Prodrug PYRRO-C3D Enhances Azithromycin Susceptibility of Nontypeable Haemophilus influenzae Biofilms. Antimicrobial Agents and Chemotherapy, 2017, 61, .	3.2	26
28	Current and future therapies for Pseudomonas aeruginosa infection in patients with cystic fibrosis. FEMS Microbiology Letters, 2017, 364, .	1.8	85
29	Qualitative Assessment of the Symptoms and Impact of Pancreatic Exocrine Insufficiency (PEI) to Inform the Development of a Patient-Reported Outcome (PRO) Instrument. Patient, 2017, 10, 615-628.	2.7	31
30	Low-Dose Nitric Oxide as Targeted Anti-biofilm Adjunctive Therapy to Treat Chronic Pseudomonas aeruginosa Infection in Cystic Fibrosis. Molecular Therapy, 2017, 25, 2104-2116.	8.2	149
31	Contributions of Composition and Interactions to Bacterial Respiration Are Reliant on the Phylogenetic Similarity of the Measured Community. Microbial Ecology, 2017, 74, 757-760.	2.8	4
32	Individualised field testing is a useful tool to evaluate difficult asthma. , 2017, , .		1
33	Primary ciliary dyskinesia exhibits dysregulated epithelial responses to non-typeable Haemophilus influenzae. , 2017, , .		0
34	Pseudomonas aeruginosainfection in cystic fibrosis: pathophysiological mechanisms and therapeutic approaches. Expert Review of Respiratory Medicine, 2016, 10, 685-697.	2.5	114
35	Cystic Fibrosis Nutrition â€“ Chewing the Fat. Paediatric Respiratory Reviews, 2016, 17, 42-44.	1.8	1
36	Lung function and nutritional status in children with cystic fibrosis and primary ciliary dyskinesia. , 2016, , .		1

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37	166 Improvements in inhalational treatment amongst children using the Philips I-neb insight online software. Journal of Cystic Fibrosis, 2015, 14, S100.	0.7	0
38	Ciprofloxacin during upper respiratory tract infections to reduce Pseudomonas aeruginosa infection in paediatric cystic fibrosis: a pilot study. Therapeutic Advances in Respiratory Disease, 2015, 9, 272-280.	2.6	3
39	Nutritional outcomes in cystic fibrosis â€“ are we doing enough?. Paediatric Respiratory Reviews, 2015, 16, 31-34.	1.8	7
40	Routine use of daily oral vitamin K to treat infants with cystic fibrosis. Paediatric Respiratory Reviews, 2015, 16, 22-24.	1.8	5
41	G385 RATNO - Reducing Antibiotic Tolerance using Nitric Oxide in Cystic Fibrosis: report of a proof of concept clinical trial. Archives of Disease in Childhood, 2014, 99, A159-A159.	1.9	10
42	Exhaled nitric oxide monitoring does not reduce exacerbation frequency or inhaled corticosteroid dose in paediatric asthma: a randomised controlled trial. Clinical Respiratory Journal, 2013, 7, 204-213.	1.6	36
43	Impact of antibiotic treatment for pulmonary exacerbations on bacterial diversity in cystic fibrosis. Journal of Cystic Fibrosis, 2013, 12, 22-28.	0.7	50
44	Children must be protected from the tobacco industry's marketing tactics. BMJ, The, 2013, 347, f7358-f7358.	6.0	2
45	A Quantitative Analysis of Ureaplasma urealyticum and Ureaplasma parvum Compared with Host Immune Response in Preterm Neonates at Risk of Developing Bronchopulmonary Dysplasia. Journal of Clinical Microbiology, 2012, 50, 909-914.	3.9	16
46	A Population-Based Study of Fish Allergy in the Philippines, Singapore and Thailand. International Archives of Allergy and Immunology, 2012, 159, 384-390.	2.1	54
47	Primary Tracheomalacia and Persistent Wheezing in Cystic Fibrosis During Infancy. Journal of Bronchology and Interventional Pulmonology, 2011, 18, 161-163.	1.4	2
48	A 2-year post-authorization safety study of high-strength pancreatic enzyme replacement therapy (pancreatin 40,000) in cystic fibrosis. Expert Opinion on Drug Safety, 2011, 10, 197-203.	2.4	11
49	Review article: enzyme supplementation in cystic fibrosis, chronic pancreatitis, pancreatic and periampullary cancer. Alimentary Pharmacology and Therapeutics, 2010, 32, 1-25.	3.7	52
50	The use of culture-independent tools to characterize bacteria in endo-tracheal aspirates from pre-term infants at risk of bronchopulmonary dysplasia. Journal of Perinatal Medicine, 2010, 38, 333-7.	1.4	22
51	Manifesting carriage of a Duchenne muscular dystrophy mutation: an unusual cause of impaired lung function in CF. Journal of the Royal Society of Medicine, 2010, 103, 27-29.	2.0	0
52	Molecular Microbiological Characterization of Preterm Neonates at Risk of Bronchopulmonary Dysplasia. Pediatric Research, 2010, 67, 412-418.	2.3	55
53	Studying bacteria in respiratory specimens by using conventional and molecular microbiological approaches. BMC Pulmonary Medicine, 2009, 9, 14.	2.0	55
54	Acute intestinal obstruction as a presentation of cystic fibrosis in infancy. Journal of Cystic Fibrosis, 2008, 7, 277-279.	0.7	21

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55	British Guideline on the Management of Asthma. Thorax, 2008, 63, iv1-iv121.	5.6	655
56	Colloidal silver for lung disease in cystic fibrosis. Journal of the Royal Society of Medicine, 2008, 101, 51-52.	2.0	8
57	Cystic Fibrosis Diagnosed After 2 Months of Age Leads to Worse Outcomes and Requires More Therapy. Pediatrics, 2007, 119, 19-28.	2.1	110
58	The Asthma Epidemic. New England Journal of Medicine, 2006, 355, 2226-2235.	27.0	1,432
59	Juvenile Laryngeal Papillomatosis. Primary Care Respiratory Journal: Journal of the General Practice Airways Group, 2006, 15, 125-127.	2.3	16
60	Idiopathic Bilateral Vocal Cord Palsy in Extremely Premature Neonates. Journal of Bronchology, 2006, 13, 221-222.	0.2	0
61	Chest x ray and high-resolution computed tomography in cystic fibrosis. Archives of Disease in Childhood, 2006, 91, 1043-1043.	1.9	0
62	Use of 16S rRNA Gene Profiling by Terminal Restriction Fragment Length Polymorphism Analysis To Compare Bacterial Communities in Sputum and Mouthwash Samples from Patients with Cystic Fibrosis. Journal of Clinical Microbiology, 2006, 44, 2601-2604.	3.9	129
63	Resolution of peanut allergy following bone marrow transplantation for primary immunodeficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2005, 60, 536-537.	5.7	25
64	The prevalence of stress urinary incontinence in patients with cystic fibrosis: an under-recognized problem. Journal of Pediatric Urology, 2005, 1, 5-9.	1.1	15
65	Maldigestion and malabsorption of 13C labelled tripalmitin in gastrostomy-fed patients with cystic fibrosis. Clinical Nutrition, 2004, 23, 347-353.	5.0	15
66	A rare cause of upper airway obstruction in a 5-year-old girl: a laryngeal web. Paediatric Anaesthesia, 2003, 13, 722-724.	1.1	15
67	Clinical application of direct sputum sensitivity testing in a severe infective exacerbation of cystic fibrosis. Pediatric Pulmonology, 2003, 35, 463-466.	2.0	9
68	Long term results of lung resection in cystic fibrosis patients with localised lung disease. Archives of Disease in Childhood, 2002, 86, 66-66.	1.9	11
69	Falsely elevated serum tobraycin levels in a patient receiving nebulised tobramycin. Journal of Cystic Fibrosis, 2002, 1, 146-147.	0.7	5
70	Editorial. Indian Journal of Pediatrics, 2000, 67, 121-122.	0.8	0
71	Somatizing disorders affecting the respiratory tract. Indian Journal of Pediatrics, 2000, 67, 129-131.	0.8	0
72	Bronchoalveolar lavage. Paediatric Respiratory Reviews, 2000, 1, 52-56.	1.8	30

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73	Faecal Elastase 1: A Marker of Exocrine Pancreatic Insufficiency in Cystic Fibrosis. Annals of Clinical Biochemistry, 1999, 36, 739-742.	1.6	30
74	rhDNase in cystic fibrosis. Thorax, 1999, 54, 750-750.	5.6	1
75	A randomised controlled trial to assess the relative benefits of large volume spacers and nebulisers to treat acute asthma in hospital. Archives of Disease in Childhood, 1999, 80, 421-423.	1.9	52
76	Interferon alpha Âtreatment of molluscum contagiosum in immunodeficiency. Archives of Disease in Childhood, 1999, 80, 77-79.	1.9	43
77	Diagnosis of cystic fibrosis : Indian perspective. Indian Journal of Pediatrics, 1999, 66, 923-928.	0.8	4
78	Pancreatic enzymes and fibrosing colonopathy. Lancet, The, 1999, 354, 249-250.	13.7	0
79	Pancreatic enzymes and fibrosing colonopathy. Lancet, The, 1999, 354, 249.	13.7	1
80	Difficult/therapyâ€resistant asthmaThe need for an integrated approach to define clinical phenotypes, evaluate risk factors, understand pathophysiology and find novel therapies. European Respiratory Journal, 1999, 13, 1198.	6.7	313
81	Colonic wall thickening is related to age and not dose of high strength pancreatin microspheres in children with cystic fibrosis. European Journal of Gastroenterology and Hepatology, 1999, 11, 181-184.	1.6	21
82	Lung resection for the treatment of severe localised bronchiectasis in cystic fibrosis patients. Acta Chirurgica Hungarica, 1999, 38, 23-5.	0.0	2
83	Diffuse Microcystic Pancreatic Enlargement in a Cystic Fibrosis Patient Causing Severe Gastrointestinal Symptoms and Successfully Treated by Total Pancreatectomy. Journal of Pediatric Gastroenterology and Nutrition, 1998, 26, 454-457.	1.8	1
84	Dietary fibre and the occurrence of gut symptoms in cystic fibrosis. Archives of Disease in Childhood, 1997, 76, 35-37.	1.9	42
85	Inflammatory pseudotumor of the trachea in a ten-month-old infant. , 1997, 23, 307-309.		9
86	Veno-venous haemodiafiltration in meningococcal septicaemia. Lancet, The, 1996, 347, 611-614.	13.7	2
87	Use of a reformulated Oka strain varicella vaccine (SmithKline Beecham Biologicals/Oka) in healthy children. European Journal of Pediatrics, 1996, 155, 706-711.	2.7	29
88	Flexible fibre-optic bronchoscopy in the management of lung complications in cystic fibrosis. Acta Paediatrica, International Journal of Paediatrics, 1996, 85, 675-678.	1.5	7
89	Bronchoscopic appearances of congenital lobar emphysema. , 1996, 21, 195-197.		29
90	Lung resection in cystic fibrosis patients with localised pulmonary disease.. Archives of Disease in Childhood, 1996, 74, 449-451.	1.9	28

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91	Use of a reformulated Oka strain varicella vaccine (SmithKline Beecham Biologicals/Oka) in healthy children. European Journal of Pediatrics, 1996, 155, 706-711.	2.7	1
92	Lung function reference values in Singaporean children aged 6-18 years.. Thorax, 1994, 49, 901-905.	5.6	27
93	Prednisolone and salbutamol in the hospital treatment of acute asthma.. Archives of Disease in Childhood, 1994, 70, 170-173.	1.9	55
94	Day care and asthma morbidity. Journal of Paediatrics and Child Health, 1994, 30, 257-259.	0.8	1
95	Effects of an acaricide on asthmatic children with house dust mite allergy. Pediatrics International, 1994, 36, 669-672.	0.5	3
96	Treating childhood asthma in Singapore: when West meets East. BMJ: British Medical Journal, 1994, 308, 1282-1284.	2.3	17
97	Use of pulse oximetry in the hospital management of acute asthma in childhood. Pediatric Pulmonology, 1993, 15, 345-349.	2.0	36
98	Audit strategies to reduce hospital admissions for acute asthma.. Archives of Disease in Childhood, 1993, 69, 202-205.	1.9	20
99	Prolonged hypoxaemia after nebulised salbutamol.. Thorax, 1993, 48, 574-575.	5.6	20
100	Prevention of viral induced asthma attacks using inhaled budesonide.. Archives of Disease in Childhood, 1993, 68, 85-87.	1.9	135
101	Use of budesonide in severe asthmatics aged 1-3 years.. Archives of Disease in Childhood, 1993, 69, 351-355.	1.9	89
102	Inhaled budesonide and behavioural disturbances. Lancet, The, 1991, 338, 634-635.	13.7	40
103	Acute wheezy bronchitis--lumping and splitting.. Archives of Disease in Childhood, 1991, 66, 751-752.	1.9	1